AMENDMENTS TO THE CLAIMS

1-2. (canceled)

- 3. (previously amended): The method of claim 26, wherein at least 90 wt% of the solvent mixture in said composition is the hydrophobic solvent.
- 4. (previously amended): The method of claim 26, wherein the hydrophobic solvent in said composition has a solubility in water of less than 0.1 wt%.
- 5. (previously amended): The method of claim 26, wherein the beneficial agent in said composition has a concentration from 0.1 mg/ml to 500 mg/ml.
- 6. (previously amended): The method of claim 26, wherein the beneficial agent in said composition has a concentration from 10 mg/ml to 500 mg/ml.
 - 7. (canceled)
- 8. (previously amended): The method of claim 26, wherein the viscosity of said composition is less than or equal to 2000 centipoise.
- 9. (previously amended): The method of claim 26, wherein less than 25% of the beneficial agent in said composition is released in 24 hours following administration *in vivo*.
 - 10. (canceled)
- 11. (previously amended): The method of claim 26, wherein the hydrophilic solvent, the hydrophobic solvent, the bioerodible polymer, and the beneficial agent in said composition form a suspension.

12. (previously amended): The method of claim 26, wherein the hydrophilic solvent, the hydrophobic solvent, the bioerodible polymer, and the beneficial agent in said composition form a solution.

13. (canceled)

- 14. (previously amended): The method of claim 30, wherein at least 55 wt% of the solvent mixture in said composition is the hydrophobic solvent.
- 15. (previously amended): The method of claim 30, wherein at least 90 wt% of the solvent mixture in said composition is the hydrophobic solvent.
- 16. (previously amended): The method of claim 30, wherein the hydrophobic solvent of said composition has a solubility in water of less than 0.1 wt%.

17-18. (canceled)

- 19. (previously amended): The method of claim 30, wherein the viscosity of said composition is less than 500 centipoise.
- 20. (previously amended): The method of claim 30, wherein the hydrophobic solvent is benzyl benzoate, the hydrophilic solvent is benzyl alcohol, the bioerodible polymer is polylactide, and the beneficial agent is a peptide or protein,
- 21. (previously amended): The method of claim 30, wherein the hydrophilic solvent, the hydrophobic solvent, the bioerodible polymer, and the beneficial agent form a solution.
- 22. (previously amended): The method of claim 30, wherein the hydrophilic solvent, the hydrophobic solvent, the bioerodible polymer, and the beneficial agent form a suspension.

23-25. (canceled)

26. (currently amended): A method of administering a beneficial agent, comprising injecting a composition comprising:

a solvent mixture comprising one or more hydrophobic solvents, wherein the total amount of hydrophobic solvent or solvents have a solubility in water of less than 1 wt% and wherein at least 55 wt% of the solvent mixture is the hydrophobic solvent and a hydrophilic solvent;

a bioerodible polymer; and

a beneficial agent,

the composition forming a solution, suspension, or gel;

into an organism through a needle, wherein the needle is no greater than a 25-gauge needle.

27. (currently amended): The method of claim 26, A method of administering a beneficial agent, comprising injecting a composition comprising:

a solvent mixture comprising one or more hydrophobic solvents, wherein the total amount of hydrophobic solvent or solvents have a solubility in water of less than 1 wt% and wherein at least 55 wt% of the solvent mixture is the hydrophobic solvent and a hydrophilic solvent;

a bioerodible polymer; and

a beneficial agent,

the composition forming a solution, suspension, or gel;

into an organism through a needle, wherein the needle is a 28-gauge needle.

28. (currently amended): The method of claim 26, A method of administering a beneficial agent, comprising injecting a composition comprising:

a solvent mixture comprising one or more hydrophobic solvents, wherein the total amount of hydrophobic solvent or solvents have a solubility in water of less than 1 wt% and wherein at least 55 wt% of the solvent mixture is the hydrophobic solvent and a hydrophilic solvent;

a bioerodible polymer; and

a beneficial agent,

the composition forming a solution, suspension, or gel;

into an organism through a needle, wherein the needle is a 30-gauge needle.

- 29. (canceled)
- 30. (currently amended): A method of administering a beneficial agent, comprising injecting a composition comprising:
 - a solvent mixture, comprising
- a hydrophobic solvent, wherein said hydrophobic solvent has a solubility in water of less than 1 wt%; and
 - a hydrophilic solvent;
 - a bioerodible polymer; and
 - a beneficial agent,

the composition forming a solution, suspension, or gel; and

wherein the viscosity of the composition is less than or equal to 2000 centipoise;

into an organism through a needle, wherein the needle is no greater than a 25-gauge needle.

- 31. (currently amended): The method of claim 30, A method of administering a beneficial agent, comprising injecting a composition comprising:
 - a solvent mixture, comprising
- a hydrophobic solvent, wherein said hydrophobic solvent has a solubility in water of less than 1 wt%; and
 - a hydrophilic solvent;
 - a bioerodible polymer; and
 - a beneficial agent,

the composition forming a solution, suspension, or gel; and

wherein the viscosity of the composition is less than or equal to 2000 centipoise;

into an organism through a needle, wherein the needle is a 28-gauge needle.

32. (currently amended): The method of claim 30, A method of administering a beneficial agent, comprising injecting a composition comprising:

a solvent mixture, comprising

<u>a hydrophobic solvent, wherein said hydrophobic solvent has a solubility in water of less</u> than 1 wt%; and

a hydrophilic solvent;

a bioerodible polymer; and

a beneficial agent,

the composition forming a solution, suspension, or gel; and

wherein the viscosity of the composition is less than or equal to 2000 centipoise;

into an organism through a needle, wherein the needle is a 30-gauge needle.

33-43. (canceled)

44. (previously amended): The method of claim 26, wherein the viscosity of said composition is less than 1000 centipoise.

45-47. (canceled)

48. (previously amended): The method of claim 30, wherein the viscosity of said composition is less than 1000 centipoise.

49-54. (canceled)

- 55. (New): The method of claim 27, wherein at least 90 wt% of the solvent mixture in said composition is the hydrophobic solvent.
- 56. (New): The method of claim 27, wherein the hydrophobic solvent in said composition has a solubility in water of less than 0.1 wt%.

57. (New): The method of claim 27, wherein the beneficial agent in said composition has a concentration from 0.1 mg/ml to 500 mg/ml.

- 58. (New): The method of claim 27, wherein the beneficial agent in said composition has a concentration from 10 mg/ml to 500 mg/ml.
- 59. (New): The method of claim 27, wherein the viscosity of said composition is less than or equal to 2000 centipoise.
- 60. (New): The method of claim 27, wherein less than 25% of the beneficial agent in said composition is released in 24 hours following administration *in vivo*.
- 61. (New): The method of claim 27, wherein the hydrophilic solvent, the hydrophobic solvent, the bioerodible polymer, and the beneficial agent in said composition form a suspension.
- 62. (New): The method of claim 27, wherein the hydrophilic solvent, the hydrophobic solvent, the bioerodible polymer, and the beneficial agent in said composition form a solution.
- 63. (New): The method of claim 27, wherein the viscosity of said composition is less than 1000 centipoise.
- 64. (New): The method of claim 28, wherein at least 90 wt% of the solvent mixture in said composition is the hydrophobic solvent.
- 65. (New): The method of claim 28, wherein the hydrophobic solvent in said composition has a solubility in water of less than 0.1 wt%.
- 66. (New): The method of claim 28, wherein the beneficial agent in said composition has a concentration from 0.1 mg/ml to 500 mg/ml.

67. (New): The method of claim 28, wherein the beneficial agent in said composition has a concentration from 10 mg/ml to 500 mg/ml.

- 68. (New): The method of claim 28, wherein the viscosity of said composition is less than or equal to 2000 centipoise.
- 69. (New): The method of claim 28, wherein less than 25% of the beneficial agent in said composition is released in 24 hours following administration *in vivo*.
- 70. (New): The method of claim 28, wherein the hydrophilic solvent, the hydrophobic solvent, the bioerodible polymer, and the beneficial agent in said composition form a suspension.
- 71. (New): The method of claim 28, wherein the hydrophilic solvent, the hydrophobic solvent, the bioerodible polymer, and the beneficial agent in said composition form a solution.
- 72. (New): The method of claim 28, wherein the viscosity of said composition is less than 1000 centipoise.
- 73. (New): The method of claim 31, wherein at least 55 wt% of the solvent mixture in said composition is the hydrophobic solvent.
- 74. (New): The method of claim 31, wherein at least 90 wt% of the solvent mixture in said composition is the hydrophobic solvent.
- 75. (New): The method of claim 31, wherein the hydrophobic solvent of said composition has a solubility in water of less than 0.1 wt%.
- 76. (New): The method of claim 31, wherein the viscosity of said composition is less than 500 centipoise.

77. (New): The method of claim 31, wherein the hydrophobic solvent is benzyl benzoate, the hydrophilic solvent is benzyl alcohol, the bioerodible polymer is polylactide, and the beneficial agent is a peptide or protein,

- 78. (New): The method of claim 31, wherein the hydrophilic solvent, the hydrophobic solvent, the bioerodible polymer, and the beneficial agent form a solution.
- 79. (New): The method of claim 31, wherein the hydrophilic solvent, the hydrophobic solvent, the bioerodible polymer, and the beneficial agent form a suspension.
- 80. (New): The method of claim 31, wherein the viscosity of said composition is less than 1000 centipoise.
- 81. (New): (New): The method of claim 32, wherein at least 55 wt% of the solvent mixture in said composition is the hydrophobic solvent.
- 82. (New): The method of claim 32, wherein at least 90 wt% of the solvent mixture in said composition is the hydrophobic solvent.
- 83. (New): The method of claim 32, wherein the hydrophobic solvent of said composition has a solubility in water of less than 0.1 wt%.
- 84. (New): The method of claim 32, wherein the viscosity of said composition is less than 500 centipoise.
- 85. (New): The method of claim 32, wherein the hydrophobic solvent is benzyl benzoate, the hydrophilic solvent is benzyl alcohol, the bioerodible polymer is polylactide, and the beneficial agent is a peptide or protein,

86. (New): The method of claim 32, wherein the hydrophilic solvent, the hydrophobic solvent, the bioerodible polymer, and the beneficial agent form a solution.

- 87. (New): The method of claim 32, wherein the hydrophilic solvent, the hydrophobic solvent, the bioerodible polymer, and the beneficial agent form a suspension.
- 88. (New): The method of claim 32, wherein the viscosity of said composition is less than 1000 centipoise.